Attachment 4.1 A

DEPARTMENT OF HEALTH AND HUMAN SERVICES Public Health Service Food and Drug Administration	APPLICATION FROM 21 CFR LASER LIGHT OR	1040.11(c)	FOR A	Form Approved, OMB N Expresion Date: Octobe See Page 4 for OMB St DOCKET NUMBER	or 31, 2000
MOTE: No laser light show, projection system, or devi-	ce may vary from compli		2 12 1 1 1 1 1 1	design or use without	the approval of
Check all applicable boxes and type or print the requested information. Submit an original and four (4) copies,	Drug /	our application to the	1-23, 12420 (igement Branch (MFA-3) Parklawn Drive, Rockvill	
1. NAME OF COMPANY MOC Shone Loser Art 2. ADDRESS OF COMPANY INCHASE ZIP CODUITY P.O.	Production	ns			
3. NAME AND TITLE OF RESPONSIBLE PERSON	Mon Heigh	LEPHONE NO. MAC	DiS	S PATE OF SL	
G. THE APPLICANT REQUESTS THE VARIANCE TO BE general the Agency will approve a variance for only the	E IN EFFECT FOR A PERI	347 520 od of	YEARS	FROM THE DATE OF I	SSUE. Un
7	PRODUCT DESCRIPT	ON AND USE		7 1.22 100 00 pay. 07 01.	
4. UST NAME AND/OR MODEL NUMBER(S) FOR THE	(E)WOH? THOU REEAL	AND PROJECTOR	s Baow	s by TaoSha	ne Laser
B. PRODUCT FOR WHICH A VARIANCE IS REQUESTE	<u> </u>	PRODUCT IS INTEN	DED TO BE U	TO TELEDIO	<u>Tecufaci</u> u:
A laser display device	\\	I. PROPUCT IS INTENDED TO BE USED AT ANY ONE LOCATION UND			
A projector for a laser light show		More than 51		han 15 days	y enneda
A laser light show		🙎 Less than 6 d	_	r.o	zanski. elina. Wl
Other (Specify)	9.	FOUR IS INTENDED			
c. [] PROJECTORS ARE INTENDED FOR SALE, LEASI OTHER LASER LIGHT SHOW PRODUCERS	E, DH LOAN 10	☐ More than 6 :	months		or previou
d. PRODUCT IS INTENDED FOR USE IN A	į	Less than one	dinom e		certrijer,
Planetarium or other dome projection structul	7e	Not applicable			ariance
Theater		Other (Specif			Accession 89A05
Hotel/motel ballroom or meeting room	n.	``		ang laser effects	
☐ Store displays ☐ Trace show or convention ☐ ☐ 75 1/5 1/5	ll remarally	Front screen			ļ
	in school	Holographic d	•		\$
D Pavilion Symnasiu		Multiple reflec		in offects	İ
	c in all			cludes scanning any a	ccessible
Outdoor arena types of	' faculitude.	uncontrolled i	ereas)		
Any quid			-	mirrors or mirrored	į
Outdoor unenclosed area USE OHIT	screage "doi	en -sudaces (Bea			
	oran ando			ating mirror balls, etc. ting mirror balls, etc.	
At only one (Freed) location	iranace.	D Fiber opticipe		ting meror balls, atc.	
At a variety of (Tour) locations	£	Foo smoke	on ones ecells	uing enhancement eff	ects
Other (Specify) Various Single	performandes	Other (Specif			
a. to be booked in the fu	TUT ASER RADIATIO	N LEVELS			
LASER MEDIUM LAT. He-Ne. etc.)	WAVE LENGTH	S (nm)		PEAK POWER (WATE	si
l HeNe			1 1	O milliwat	T< 1
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			2 - 25 - 5 - 5 - 5	The second secon	
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9. IF ANY LASER RADIATION IS PULSED OR SCANNED SCANNING PULSED OR SCANNED	WE THE PULSE DURG	TION AND BATE AL	ND SCANNING	FREQUENCY AND AN	APUTUDE
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which Urely on scani	ning to me	בו ננננע	7, CXP	10 701 € 101	1
10. REASON FOR REQUESTING VARIANCE					
Compliance with the limits of 21 CFR 1040.1	T(c) would restrict the li	ntended use of the	product becau	se compliance would	
inthit the output power to the extent that the	desued effects would no	t be sufficiently wai	pie	A	IACY
Other or additional explanation (Specify)				L	2025
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ATTachment 4.1B

It is proposed to deviate from the provisions of 21 CFR 1040.11(c) in that the accessible emission level would exceed the accessible emission limits specified in 21 CFR 1040.11(c). It is proposed to deviate from the provisions of 21 CFR 1040.11(c).
12. ADVANTAGES TO BE DERIVED FROM SUCH DEVIATION Lazer light shows and displays are accepted popular modic in entertainment and the arts. Use of power levels in excess of the limits imposed by 21 CFR 1040.11(c) is necessary to acrees the required effects in these media.
Additional advantage is that additional power will enhance tests in locations which cannot be fully darkened
13. EXPLAIN THE ALTERNATE MEANS OF RADIATION PROTECTION TO BE PROVIDED. (Check as many dokes as apply. In item 14 'Romarks,' justify any oders not checked, using additional shoots as necessary. State any other means of radiation protection that will be used.)
a. All laser products, systems, shows, and projectors will be certified to comply with 21 CFR 1040.10 and the conditions of this variance and will be reported as required by 21 CFR 1002.10 AND 1002.11 using the reporting guides provided for such purpose. These actions will be accomplished prior to any introduction into commerce.
b. A Effects not specifically indicated in this variance application will not be performed. No other effects will be added until an amendment to the variance has been obtained and the required reports or supplements, as applicable, have been submitted.
CAS Scanning, projection, or reflection of laser and collateral radiation (Light snow radiation) into audience or other accessible uncontrolled areas will not be permitted except for diffuse reflections produced by the atmosphere, added atmospheric scattering media, and target screens.
d. Laser radiation levels in excess of the limits of Class I will not be permitted at any point less than 3.0 meters above any surface upon which persons other than operators, performers, or employees are permitted to stand or 2.5 meters pelow or in lawral separation from any place where such persons are permitted to be. Operators, performers, and employees will not be required or allowed to view radiation above the limits of Class I or be exposed to radiation above the limits specified in 21 CFR 1040.11(c).
e. Any product which relies on scanning to meet access, exposure, or product class limits will incorporate a scanning safeguard system which directly senses scanner motion and which will react fast enough to preclude exceeding the applicable limit.
f. All laser light shows shall be under the direct and personal control of trained, competent operator(s). The operator(s) will:
(1) Be an employee of the variance holder who will be responsible for the training and the conduct of the operator:
(2) Be located where all beam pains can be directly observed at all times; and
(3) Immediately terminate the emission of light show radiation in the event of any unsale condition; or, for outdoor shows, upon request by any air traffic control officials.
g. The maximum laser projector output power will not exceed the level required to obtain the intended effects.
h. The projection system (i.e., the projector and all other components used to produce the lighting effects) will be securely mounted or immobilized to provent unintended movement or missignment. Seam marking will be provided as an interent part of the system design to provent overfilling of screens, beam stops, targets, etc.
i. Laser projectors will not be delivered to any other party under an agreement of sale, lease, or loan unless and until the recipient demonstrates that they have a variance in effect at the time projectory that points them to produce laser light shows incorporating such projector(s).
j. In addition to the requirements of 21 CFR 1040.10(h), the manufacturer of laser projectors/systems will provide to parties who purchase, lease, or borrow the equipment, adequate users' instructions for sale installation and operation which explain the responsibility the recipient as an independent light show manufacturer to submit the required reports and apply for and obtain a variance from CDRH prior to introduction into commerce of any laser Eght zhows.
The requirements of 21 CFR 1002.30(a)(1) and (2) will be eccomplished through the use of written procedures for setup, alignment, testing, and performance of each show, these procedures will be in sufficient detail to ensure compliance with 21 CFR 1040.10, the conditions of this variance, and the control of access to radiation areas using the procedures described in the ANS/2136.1 standards in the safe use of lasers (American National Standards Inspire, 1430 Broadway, New York, NY 10018) or any other equivalent usor consensus standard and, where applicable, state or local requirements. Laser radiation areas which can contain radiation levels above the limits specified in 21 CFR 1040.11(c) will be clearly identified by the posting of warning signs and/or restricting access through physical means (such as pressure switches, photo cells, parriars, guards, etc.). These requirements apply to temporary areas (such as during set up and alignment procedures) and to [and] or permanent areas. The variance holder will retain the records of these procedures and the results of all tests as required by 21 CFR 1002.31. A copy of the variance application, the approval letter, current procedures and records relating to each particular show will be with the operator or other responsible individual and will be made available for inspection by FDA and other responsible authorities.

Attachment 4.1 C

- I. Advance written notification will be made as carly as possible to appropriate taderal, state, and local authorities providing show itinerary with dates and locations clearly and completely identified, and a basic description of the proposed effects including a statement of the maximum power output intended. Such notifications will be made, but not necessarily be limited. To:
 - (1) The Center for Devices and Radiological Health. Office of Compliance (HFZ-342), 2098 Gaither Road, Rockville, MD 20850, providing the initial and closing dates for fixed installations and the itinerary for mobile shows. In addition, unless all aspects of each show have been reported and accession numbers clearly referenced, each notice will include detailed descriptions of each show and a listing of all effects to be performed in sufficient detail to confirm compliance with the regulations and this variance.
 - (2) The Federal Aviation Administration (FAA) for any projections into open airspace at any time (i.e., including set up, dignment rehearsels, performances, etc.). If the FAA objects to any laser effects, the objections will be resolved and any conditions requested by FAA will be adhered to. If these conditions cannot be may the objectionable effects will be deleted from the show
 - (3) State and local radiation control offices/agencies for all shows to be performed within their jurisdictions. All requirements of state and local law will be satisfied and any objections raised by local authorities will be resolved or the effects deleted. (A list of federal and state offices is available from the Center for Devices and Radiological Health upon request.)

14. REMARKS

No effects will be performed which rely on scanning to meet access, exposure, or product J class limits

13 j No projectors/systems will be sold, leased, or loaned to any other parties.

CERTIFICATION

I CERTIFY that all of the above information and statements are true, complete, and correct to the best of my knowledge and acknowledge that my variance application may be denied or my variance may be revoked if this application is found to be false, misleading or incorrect in any material way. I have submitted and will support required by 21 CFR 1002-10 and 1002-11 on the laser equipment and show(s). I further understand that I may be required by regulation or by the Director. Center for Devices and Radiological Health, to supply such other information as may be necessary to available and act on this application.

15. SIGNATURE Abyanshi

Rozanski Kenneth Rozanski to

17. TITLE Technician